AMENDMENTS TO THE CLAIMS

- 110. (Currently Amended) A method of preparing [[a]] xenotransplantable porcine <u>pancreatic</u> islet <u>cells</u> comprising the steps of:
- (i) harvesting the pancreas of piglets from a piglet, the piglet having an age of between -20 to +10 days relative to full term gestation, [[and]]
 - (ii) exposing pancreatic islet cells to nicotinamide, and
- [[(ii)]] (iii) extracting the pancreatic [[β]] islet cells from the harvested pancreas; and (iii) exposing the islets to nicotinamide either before or after either the steps of harvesting or extracting; the method resulting in a xenotransplantable islet cell.
- 111. (Currently Amended) The method as claimed in claim 110 wherein the piglets are from -7 to +10 days full term gestation the piglet has an age of between -7 and +10 days relative to full term gestation.
- 112. (Currently Amended) The method as claimed in claim 110 wherein the step of **xtraction* extraction* includes the use of human Liberase.
- 113. (Currently Amended) The method of claim 110 wherein the harvested pancreas is <u>bathed in</u> a <u>mammalian albumin solution</u> in a <u>supportive mammalian albumin</u> substantially free of non-human microbiological agents.
- 114. (Previously Presented) The method of claim 113 wherein the mammalian albumin comprises human serum albumin (HSA).

- 115. (Cancelled) The method of claim 110 wherein the step of exposing occurs after the step of extracting.
- 116. (Currently Amended) The method as claimed in claim 110 further comprising the step of treating the islets with a compound selected from the group consisting of: Insulin-Like Growth Factor 1 (IGF-1) one of IGF-1 and the N-terminal tripeptide of IGF-1 (GPE).
- 117. (Currently Amended) The method as claimed in claim 116 wherein the step of treating the islets comprises the treating thereof with GPE compound comprises GPE.
- 118. (Currently Amended) The method as claimed in claim 116 wherein the amount of compound used to treat the islets is greater when using a piglet further from full-term gestation, and is less when using a piglet closer to full-term gestation the exposure to either of IGF-1 or GPE is greater for those cells from piglets furthest from full term gestation.
- 119. (Canceled) The method as claimed in claim 116 wherein the exposure to IGF-1 is unrelated to their relationship with full term gestation.
- 120. (Previously Presented) The method as claimed in claim 110 further comprising the step of subjecting at least one of the pancreas and the islets to a trauma protecting agent.
- 121. (Previously Presented) The method as claimed in claim 120 wherein the trauma protecting agent comprises an anesthetic agent.

- 122. (Previously Presented) The method as claimed in claim 121 wherein the anaesthetic agent comprises lignocaine.
- 123. (Previously Presented) The method as claimed in claim 110 further comprising the step of mechanically reducing the harvested pancreas in the presence of an islet trauma protecting agent.
- 124. (Currently Amended) The method as claimed in claim 110 further comprising the step of associating a quinaline contacting a quinalone antibiotic with the islet cells.
- 125. (Currently Amended) The method as claimed in claim 124 wherein the quinaline quinalone antibiotic comprises ciproxin.
- 126. (Previously Presented) The method as claimed in claim 110 further comprising the steps of encapsulating the islet cells with a biocompatible xenotransplantable material, said material being both glucose and insulin porous, the biocompatible xenotransplantable material comprising a suitable alginate in ultra pure form.
- 127. (Previously Presented) The method as claimed in claim 126 wherein the step of encapsulating comprises the steps of:
- (i) presenting islets and the suitable alginate in ultra pure form into a source of compatible cations; and
 - (ii) entrapping the islets in a cation-alginate gel.

- 128. (Previously Presented) The method as claimed in claim 127 wherein the cation alginate gel comprises calcium-alginate gel.
- 129. (Previously Presented) The method as claimed in claim 128 wherein the alginate in ultra pure form comprises sodium alginate.
- 130. (Previously Presented) The method as claimed in claim 129 wherein a resulting solution of islet and sodium alginate is of 1.6% w/w.
- 131. (Previously Presented) The method as claimed in claim 129 wherein the suitable cation comprises calcium chloride.
- 132. (Previously Presented) The method as claimed in claim 127 further comprising the steps of:
 - (i) coating the gel encased islets with a positively charged material; and
 - (ii) providing an outer coat of a suitable alginate.
- 133. (Previously Presented) The method as claimed in claim 132 wherein the positively charged material comprises a poly-L-ornithine.
- 134. (Previously Presented) The method as claimed in claim 132 further comprising the step of liquefying the gel entrapping the islets.

- 135. (Previously Presented) The method as claimed in claim 134 wherein the step of liquefying comprises the step of exposing the gel to sodium citrate.
- 136. (Previously Presented) The method as claimed in claim 134 further comprising the steps of:
 - (i) washing the outer coat of a suitable alginate; and
 - (ii) recoating the outer coat with a suitable alginate.
- 137. (Previously Presented) The method as claimed in claim 126 wherein the step of encapsulation produces at least one capsule.
- 138. (Previously Presented) The method as claimed in claim 137 wherein the at least one capsule includes a plurality of islet cells.
- 139. (Previously Presented) The method as claimed in claim 138 wherein the at least one capsule includes at least three islet cells.
- 140. (Previously Presented) The method as claimed in claim 137 wherein the at least one capsule includes a diameter of about 300 to 400 microns.
- 141. (Previously Presented) A method of treating a mammalian patient suffering from diabetes, the method comprising the steps of:
 - (i) extracting pancreatic β islet cells from the harvested pancreas; and
 - (ii) encapsulating the islet cells with a biocompatible xenotransplantable material, said

material being both glucose and insulin porous;

- (iii) introducing a trauma protecting agent during or prior to the step of encapsulating; and
- (iv) transplanting into the mammalian patient an effective amount of viable islet cells capable of producing insulin in the patient.
- 142. (Previously Presented) The method as claimed in claim 141 wherein the trauma protecting agent is selected from suitable anesthetic agents.
- 143. (Previously Presented) The method as claimed in claim 142 wherein the trauma protecting agent comprises lignocaine.
- 144. (Previously Presented) The method as claimed in claim 141 further comprising the step of subjecting the patient to a cholesterol lowering drug regime prior to, during or after the step of transplanting.
- 145. (Previously Presented) The method as claimed in claim 144 wherein the drug regime comprises one of the "statin" family.
- 146. (Previously Presented) The method as claimed in claim 145 wherein the drug regime comprises one of the group consisting of pravastatin and simvistatin.

- 147. (Previously Presented) The method as claimed in claim 141 further comprising the step of prescribing to the patient, prior to or after the transplanting step, a casein-free diet.
- 148. (Previously Presented) A method of treating a mammalian patient suffering from diabetes, the method comprising the steps of:
 - (i) extracting pancreatic β islet cells from the harvested pancreas; and
- (ii) encapsulating the islet cells with a biocompatible xenotransplantable material, said material being both glucose and insulin porous;
- (iii) transplanting into the mammalian patient an effective amount of viable islet cells capable of producing insulin in the patient; and
- (iv) subjecting the patient to a cholesterol lowering drug regime prior to, during or after the step of transplanting.
- 149. (Previously Presented) The method as claimed in claim 148 wherein the drug regime comprises one of the "statin" family.
- 150. (Previously Presented) The method of claim 149 wherein the drug regime comprises one of the group consisting of pravastatin and simvistatin.
- 151. (Previously Presented) The method as claimed in claim 150 further comprising the step of prescribing to the patient, prior to or after the transplanting step, a casein-free diet.

- 152. (New) A method of preparing xenotransplantable porcine islet cells comprising the steps of:
 - (i) providing a piglet, the piglet having an age of between
- -20 and +10 days relative to full term gestation,
 - (ii) harvesting the pancreas of the piglet,
- (iii) extracting pancreatic .beta. islet cells from the harvested pancreas and simultaneously exposing the .beta. islet cells to nicotinamide; the method resulting in a xenotransplantable islet cell.
- 153. (New) A method of preparing xenotransplantable porcine islet cells comprising the steps of:
- (i) providing a piglet, the piglet having an age of between -20 and +10 days relative to full term gestation,
- (ii) harvesting the pancreas of the piglet and simultaneously exposing .beta. islet cells to nicotinamide, and
- (iii) extracting the pancreatic .beta. islet cells from the harvested pancreas; the method resulting in a xenotransplantable islet cell.
- 154. (New) The method of claim 110 wherein the piglet has not reached full term gestation.
- 155. (New) The method of claim 153 wherein the piglet has not reached full term gestation.
 - 156. (New) The method of claim 154 wherein the piglet has not reached full term gestation.